- 1. (Cancelled)
- (Cancelled)
- 3. (Previously Amended) A compound selected from the group consisting of:
- (1α, 5α, 6α)-6N-[3-benzyl-3-azabicyclo[3,1.0]hexyl]-3-methyl-4-oxo-α-phenyl-4H-1-benzopyran-8-carboxamide (Compound No. 1):
- (1α, 5α, 6α)-6N-[3-(4-cyanobenzyl)-3-azabicyclo[3.1.0]hexyl]-3-methyl-4-oxo-2-phenyl-4H-1-benzopyran-8-carboxamide (Compound No. 2);
- (1α,5α, 6α)-N-[3-benzyl-3-azabicyclo [3.1.0] hexyl-6-(aminomethyl)-yl]-3-methyl-4-oxo-2-phenyl-4H-1-benzopyran-8-carboxamide (Compound No. 3);
- (1α,5α,6α)-N-[3-(4-methyl-3-pentyl)-3-azabicyclo[3.1.0]hexyl-6-(aminomethyl)-yl]-3-methyl-4-oxo-2-phenyl-4H-1-benzopyran-8-carboxamide (Compound No. 4); and
- N-[3-benzyl-3-azabicyclo[3.1.0]hexyl-1-(aminomethyl)-yl]-3-methyl-4-oxo-2-phenyl-4H-1-benzopyran-8-carboxamide (Compound No. 5).
- (Currently Amended) A pharmaceutical composition comprising a
 pharmaceutically effective amount of a compound as defined in claim 4, 2 or 3 together
 with pharmaceutically acceptable carriers, excipients, or diluents.
- 5. (Currently Amended) A method for treatment of an animal or a human suffering from a disease or disorder of the respiratory, urinary and gastrointestinal systems, wherein the disease or disorder is urinary incontinence, lower urinary tract symptoms (LUTS), bronchial asthma, chronic obstructive pulmonary disorders (COPD), pulmonary fibrosis, irritable bowel syndrome, obesity, diabetes, and gastrointestinal hyperkinesis, comprising administering to said animal or human, a therapeutically effective amount of a compound of claim 3 having the structure of Formula I.

Formula I

or its pharmaceutically acceptable salts, pharmaceutically acceptable solvates, esters, enantiomers, diastereomers, Noxides, prodrugs, metabolites, wherein:

W represents (CH₂)_n, where p represents 0 to 1;

X represents an oxygen, sulphur, nitrogen or no atom;

Y represents CHR₁CO₂ wherein R₁ represents hydrogen or methyl or (CH₂)q wherein a represents 0 to 4:

Z—represents oxygen, sulphur, NR₂, wherein R₂ represents hydrogen, C₁₋₆ alkyl; Q—represents (CH₂)n wherein n represents 0 to 4, or CHR₃ wherein R₂ represents H, OH, C₁₋₆, alkyl, alkenyl, alkoxy or CH₂CHR₄ wherein R₄-represents H, OH, lower alkyl—(C₁-C₄) or lower alkoxy (C₁-C₄);

Rs and R6 are independently selected from COOH, H, CH3, CONH2, NH2, CH2NH2;

 R_2 represents $C_1 \cdot C_{15}$ saturated or unsaturated aliphatic hydrocarbon groups in which any 1 to 6 hydrogen atoms may be substituted with the group independently selected from halogen, arylalkyl, arylalkenyl, heteroarylalkyl or heteroarylalkenyl having 1 to 2 hetero atoms selected from a group consisting of nitrogen, oxygen and sulphur atoms with option that any 1 to 3 hydrogen atoms on the ring in said arylalkyl, arylalkenyl, hetero arylalkenyl group may be substituted with lower alkyl $(C_1 \cdot C_4)$, lower perhalo alkyl $(C_1 \cdot C_4)$, eyano, hydroxyl, nitro, lower alkoxycarbonyl, halogen, lower alkoxy $(C_1 \cdot C_4)$, lower perhaloalkoxy $(C_1 \cdot C_4)$; unsubstituent amino, N-lower alkylamino $(C_1 \cdot C_4)$, N-lower alkylamino earbonyl $(C_1 \cdot C_4)$;

Aryl rings may be unsubstituted or substituted by R_8 and R_9 in which any one to three substituents may be independently selected from lower alkyl (C_1, C_4) , trifluoromethyl, eyano, hydroxy, nitro, lower alkoxy (C_1, C_4) , amino or lower alkylamino; and R_{10} represents aryl which may be substituted with one or more substituent.

6. (Currently Amended) The method according to claim 5 for treatment of an animal or a human suffering from a disease or disorder of the respiratory, urinary and gastrointestinal systems, wherein the disease or disorder is urinary incontinence, lower urinary tract symptoms (LUTS), bronchial asthma, chronic obstructive pulmonary disorders (COPD), pulmonary fibrosis, irritable bowel syndrome, obesity, diabetes, and gastrointestinal hyperkinesis, comprising administering to said animal or human, a therapeutically effective amount of a compound of claim 3 having the structure of Formula II, and its pharmaceutically acceptable salts, pharmaceutically acceptable, esters, enantiomers, diastercomers, prodrugs, polymorphs, or metabolites, wherein Rs, Rs, Rug; Rz, WyX, Y, Z, Q are as defined for Formula I.

Formula II

7.- 8. (Previously Cancelled).

(Currently Amended) The method for treatment or prophylaxis of an animal or a
human suffering from a disease or disorder of the respiratory, urinary, and
gastrointestinal systems, wherein the disease or disorder is urinary incontinence, lower
urinary tract symptoms (LUTS), bronchial asthma, chronic obstructive pulmonary

RLL-283US Mehta et al. 10/540,062 Page 5 of 8

disorders (COPD), pulmonary fibrosis, irritable bowel syndrome, obesity, diabetes, and gastrointestinal hyperkinesis, comprising administering to said animal or human, a therapeutically effective amount of the pharmaceutical composition according to claim 4.

10. (Previously Cancelled).

11.-18. (Cancelled).